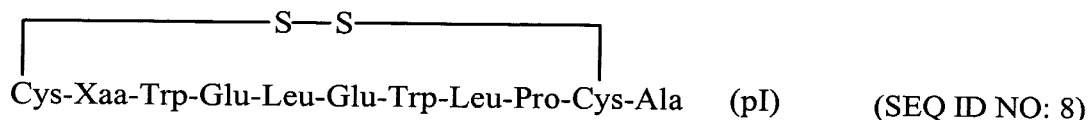


*Amendments to the Claims*

9. (Previously Amended) A cyclic peptide having the structure:



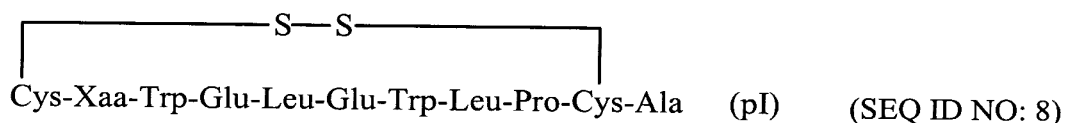
wherein Xaa is Tyr or Ala.

11. (Previously Amended) A peptide of claim 9, wherein Xaa is Tyr (SEQ ID NO:10).

12. (Previously Added) The cyclic peptide of claim 9, wherein the cyclic peptide is in contact with an aqueous solution.

20. (Currently Amended) A method of identifying or designing a phospholamban deactivator, comprising the steps of:

- (a) obtaining a three dimensional structure of a cyclic peptide



wherein Xaa is Tyr or Ala;

- (b) creating a three dimensional model of a complex of the cyclic peptide bound to phospholamban cytosolic domain or ligand-binding portion of the phospholamban cytosolic domain ~~thereof~~;

- (c) employing the three dimensional model of the complex to identify a ligand binding site on the phospholamban cytosolic domain or on the ligand-binding portion of the phospholamban cytosolic domain ~~thereof~~, wherein the ligand binding site is the site at which the phospholamban deactivator binds to the phospholamban cytosolic domain or to the ligand-binding portion of the phospholamban cytosolic domain when the phospholamban deactivator is bound to phospholamban;

D<sup>1</sup> (d) selecting a candidate molecule that ~~is capable of interacting~~  
computationally binds with the ligand binding site ~~on the phospholamban cytosolic~~  
~~domain or ligand-binding portion of the phospholamban cytosolic domain thereof~~ or  
that possesses good steric and electrostatic complementarity with the ligand binding  
site; and

(e) identifying the selected candidate molecule as a potential  
phospholamban deactivator, wherein the potential phospholamban deactivator can be  
subsequently synthesized and tested for its ability to function as the phospholamban  
deactivator.

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21. (Previously Added) The method of claim 20, which further comprises:

(f) synthesizing the potential phospholamban deactivator and testing the  
synthesized potential phospholamban deactivator for activation of CaATPase in the  
presence of phospholamban.

22. (Previously Added) The method of claim 20, wherein step (a) comprises  
obtaining a first set of atom coordinates defining the three dimensional structure of the  
cyclic peptide.

23. (Currently Amended) The method of claim 20, wherein step (b)  
comprises:

D<sup>2</sup> (i) obtaining the first set of atom coordinates defining the three  
dimensional structure of the cyclic peptide of step (a);

(ii) obtaining a second set of atom coordinates defining the  
phospholamban cytosolic domain or ligand-binding portion of the phospholamban  
cytosolic domain ~~thereof~~; and

(iii) employing a computer-aided molecular modeling program to combine  
the first set of atom coordinates with the second set of atom coordinates to create a  
three dimensional model of a complex of the cyclic peptide bound to phospholamban  
cytosolic domain or ligand-binding portion of the phospholamban cytosolic domain  
~~thereof~~.

24. (Currently Amended) The method of claim 20, wherein step (c) employs a computer-aided molecular modeling program to identify the ligand binding site on the phospholamban cytosolic domain or the ligand-binding portion of the phospholamban cytosolic domain ~~thereof~~.

25. (Currently Amended) The method of claim 20, wherein step (d) employs a computer-aided molecular modeling program to identify the compound capable of interacting with the ligand binding site of the phospholamban cytosolic domain or ligand-binding portion of the phospholamban cytosolic domain thereof.

26. (Currently Amended) The method of claim 20, wherein step (d) comprises:

D<sup>2</sup>  
(i') providing atom coordinates defining a three-dimensional structure of the phospholamban cytosolic domain or ligand-binding portion of the phospholamban cytosolic domain ~~thereof~~ that is in a conformation which allows binding of the phospholamban deactivator;

(ii') combining the atom coordinates defining the three-dimensional structure of the phospholamban cytosolic domain or ligand-binding portion of the phospholamban cytosolic domain of step (i') with a set of atom coordinates defining a three dimensional structure of a candidate molecule;

(iii') employing a computer-aided molecular modeling program, with the atom coordinates defining the three-dimensional structure of the phospholamban cytosolic domain or ligand-binding portion of the phospholamban cytosolic domain and the atom coordinates defining the three dimensional structure of the candidate molecule, to evaluate the ability of the candidate molecule to ~~interact with~~ bind to the ligand binding site of the phospholamban cytosolic domain or ligand-binding portion of the phospholamban cytosolic domain ~~thereof~~; and

(iv') selecting the candidate molecule that interacts favorably with the ligand binding site of the phospholamban cytosolic domain or ligand-binding portion of the phospholamban cytosolic domain ~~thereof~~, or that possesses good steric and electrostatic complementarity with the ligand binding site.

27. (Currently Amended) The method of claim 26, wherein the atom coordinates defining the three-dimensional structure of the phospholamban cytosolic domain or ligand-binding portion of the phospholamban cytosolic domain ~~thereof~~ are ~~derived~~ obtained from the three dimensional model of the complex created in step (b).

28. (Currently Amended) The method of claim 26, wherein step (iii') comprises:

(iiia') performing a fitting operation between the candidate molecule and the ligand binding site of the phospholamban cytosolic domain or ligand-binding portion of the phospholamban cytosolic domain ~~thereof~~; and

(iiib') analyzing the results of the fitting operation to quantify association between the candidate molecule and the ligand binding site of the phospholamban cytosolic domain or ligand-binding portion of the phospholamban cytosolic domain ~~thereof~~.

29. (Currently Amended) The method of claim 26, wherein step (iii') comprises:

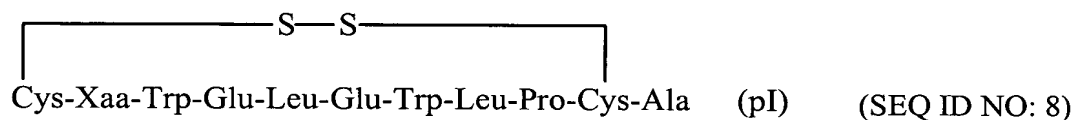
(iiia'') displaying in a graphical format a protein structure encoded by the combination of the atom coordinates defining the three-dimensional structure of the phospholamban cytosolic domain or ligand-binding portion of the phospholamban cytosolic domain and the atom coordinates defining the three dimensional structure of the candidate molecule; and

(iiib'') visually inspecting the protein structure displayed in the graphical format to evaluate the ability of the candidate molecule to ~~interact with~~ bind to the ligand binding site of the phospholamban cytosolic domain or ligand-binding portion of the phospholamban cytosolic domain ~~thereof~~.

30. (Previously Added) The method of claim 20, wherein Xaa of the cyclic peptide is Tyr (SEQ ID NO:10).

31. (Currently Amended) A method of identifying a target area on the surface of phospholamban, comprising the steps of:

- (a) obtaining a three dimensional structure of a cyclic peptide



wherein Xaa is Tyr or Ala;

- (b) creating a three dimensional model of a complex of the cyclic peptide bound to phospholamban cytosolic domain or ligand-binding portion of the phospholamban cytosolic domain thereof; and

(c) employing a computer-aided molecular modeling program and the three dimensional model of the complex to identify the target area on the surface of phospholamban, wherein said target area is the site at which a phospholamban deactivator binds to the phospholamban cytosolic domain or to the ligand-binding portion of the phospholamban cytosolic domain when said phospholamban deactivator is bound to phospholamban.

32. (Previously Added) The method of claim 31, wherein Xaa of the cyclic peptide is Tyr (SEQ ID NO:10).

33. (Previously Added) A method of preventing inhibition exerted by phospholamban on CaATPase in a cardiac cell, comprising introducing the cyclic peptide of claim 9 into a cardiac cell.